Appl. No. 10/789,807 Amdt. dated June 15, 2006 Reply to Office Action of January 3, 2006

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1. (Original) A method for differentiating monocytic dendritic cell precursors into immature dendritic cells, comprising:
- a) providing a cell population comprising non-activated monocytic dendritic cell precursors;
- b) contacting the non-activated dendritic cell precursors in a culture vessel with a dendritic cell culture media supplemented with granulocyte-macrophage colony stimulating factor in the absence of additional cytokines.
 - 2. (Canceled)
- 3. (Currently amended) The method according to claim [[2]] 1, wherein activation of the monocytic dendritic cell precursor cells is prevented by inhibiting the adhesion of the precursor cells to the culture vessel.
- 4. (Withdrawn) The method according to claim 3, wherein the adhesion of the monocytic dendritic cell precursor cells is inhibited by contacting the cells with a dendritic cell culture medium comprising a high concentration of an animal or human protein.
- 5. (Withdrawn) The method according to claim 4, wherein the animal or human protein is an albumin, serum, plasma, gelatin, or poly-amino acid.
- 6. (Withdrawn) The method according to claim 1, wherein the activation of the monocytic dendritic precursor cell is inhibited by contacting the cells with a dendritic cell culture media comprising a metal chelator.
- 7. (Withdrawn) The method according to claim 6, wherein the metal chelator comprising EDTA, or EGTA.

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- 8. (Original) The method according to claim 3, wherein the adhesion of the monocytic dendritic cell precursor to the culture vessel is inhibited by contacting the cells with a low cellular avidity culture vessel.
- 9. (Original) The method according to claim 8, wherein the low cellular avidity culture vessel comprises polypropylene, Teflon[®], or PFTE.
- 10. (Withdrawn) The method according to claim 5, wherein the protein is human serum albumin.
- 11. (Withdrawn) The method according to claim 3, wherein the human serum albumin is present at a concentration of at least 1 %.
- 12. (Withdrawn) The method according to claim 11, wherein the human serum albumin is present at a concentration of about 2 % to about 10 %.
- 13. (Original) The method according to claim 1, wherein the dendritic cell culture medium is a serum free medium.
- 14. (Original) The method according to claim 1, wherein the cell population comprises peripheral blood, a leukapheresis product, an apheresis product, cord blood, spleen, lymph node, thymus, or bone marrow.
- 15. (Original) The method according to claim 14, wherein the cell population has been cryopreserved.
- 16. (Withdrawn) The method according to claim 4, wherein the culture vessel comprises, polystyrene, glass coated polystyrene, styrene or glass.
- 17. (Original) The method according to claim 14, wherein the dendritic cell precursors are further enriched by tangential flow filtration.

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- 18. (Currently amended) The method according to claim 17, wherein the filter has a pore size of 5.5 micron, the recirculation (input) rate [[was]] is about 1400 ml/min, the filtration rate [[was]] is about 17 ml/min, and the filtration time [[was]] is about 90 min.
- 19. (Original) The method according to claim 1, further comprising contacting the differentiated dendritic cell precursors with an antigen of interest for a time period sufficient for antigen uptake.
- 20. (Original) The method according to claim 19, further comprising contacting the differentiated dendritic cell precursors with a dendritic cell maturation agent.
- 21. (Currently amended) The method according to claim 20, wherein the dendritic cell maturation agent comprises [[is]] Bacillus Calmette-Guerin (BCG), lipopolysaccharide (LPS), TNFα, Interferon gamma (IFNγ), or combinations thereof.
- 22. (Original) The method according to claim 21, wherein the maturation agent is a combination of BCG and IFNγ.
- 23. (Original) The method according to claim 19, wherein the antigen is a tumor specific antigen, a tumor associated antigen, a viral antigen, a bacterial antigen, tumor cells, a nucleic acid encoding the antigen isolated from a tumor cell, bacterial cells, recombinant cells expressing an antigen, a cell lysate, a membrane preparation, a recombinantly produced antigen, a peptide antigen, or an isolated antigen.
- 24. (Withdrawn) The method according to claim 10, further comprising cryopreservation of the dendritic cells.
- 25. (Withdrawn). The method according to claim 8, wherein the monocytic dendritic cell precursor cells are contacted with a dendritic cell culture medium comprising a high concentration of an animal or human protein.

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- 26. (Withdrawn) The method according to claim 25, wherein the animal or human protein is an albumin, serum, plasma, gelatin, or poly-amino acid.
- 27. (Withdrawn) The method according to claim 26, wherein the protein is human serum albumin.
- 28. (Withdrawn) The method according to claim 27, wherein the human serum albumin is present at a concentration of at least 1 %.
- 29. (Withdrawn) The method according to claim 27, wherein the human serum albumin is present at a concentration of about 2 % to about 10 %.